- 2 (a) one or more transfer chambers;
- 3 (b) a substrate handling member disposed in each of the one or more transfer 4 chambers;
- one or more processing chambers, each processing chamber defining at least one isolated processing region therein, wherein each processing region is connected to the one or more transfer chambers;
 - (d) one or more loadlock chambers in communication with the one or more transfer chambers; and
 - (e) one or more multi-slot substrate pre-heating modules in communication with the one or more transfer chambers.
- 1 2. The apparatus of claim 1, further comprising one or more multi-slot cooling stations disposed within the loadlock chamber.
- 1 3. The apparatus of claim 1, further comprising a vacuum pump in fluid communication with the loadlock chamber.
 - 4. The apparatus of claim 1, further comprising a vacuum pump in fluid communication with each processing region in the one or more processing chambers.
- 1 5. The apparatus of claim 1, wherein each processing chamber has two isolated 2 processing regions.
- 1 6. The apparatus of claim 1, wherein each processing region includes a gas
- distribution assembly disposed therein and each gas distribution assembly shares process
- 3 gases from one or more gas sources.
- 1 7. The apparatus of claim 1, further comprising a remote plasma system having an RF

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- 8. The apparatus of claim 1, wherein a remote plasma system is in fluid communication with each processing region.
- 9. The apparatus of claim 2, further comprising a high pressure deposition module connected to the one or more load lock chambers.
- 1 10. The apparatus of claim 9, wherein the high pressure deposition module is a spin-on dielectric module comprising one or more substrate stripping chambers.
 - 1 11. The apparatus of claim 1, wherein the one or more multi-slot pre-heating modules are disposed within the loadlock chamber.
 - 1 12. An apparatus for processing substrates, comprising:
 - 2 (a) a high pressure deposition module;
 - 3 (b) a first transfer chamber in communication with the high pressure deposition 4 module;
- 5 (c) a loadlock chamber in communication with the first transfer chamber;
- 7 (d) a second transfer chamber in communication with the loadlock chamber;
- 8 (e) a multi-slot substrate pre-heating module in communication with the first 9 transfer chamber;
- 10 (f) a substrate handling member disposed in the second transfer chamber; and
- one or more processing chambers, each processing chamber defining at
- least one isolated processing region therein, wherein each processing region is connected
- to the second transfer chamber.
- 1 13. The apparatus of claim 12, wherein the high pressure deposition module comprises:
 - (a) one or more substrate spinner chambers;

3 (d) one or more silylation deposition chambers; and

4 (e) a second substrate handling member disposed in the high pressure deposition module.

1 14. The apparatus of claim 12, further comprising one or more multi-slot cooling stations disposed within each of the one or more loadlock chambers.

5. The apparatus of claim 12, further comprising a vacuum pump in fluid communication with the one or more loadlock chambers.

1 16. The apparatus of claim 12, further comprising a vacuum pump in fluid

2 communication with each processing region.

1 17. The apparatus of claim 12, wherein each processing chamber has two isolated

2 processing regions.

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1 18. The apparatus of claim 12, wherein each processing region includes a gas

2 distribution assembly disposed therein and each gas distribution assembly shares process

3 gases from one or more gas sources.

1 19. The apparatus of claim 12, further comprising a remote plasma system having a RF

2 generator connected to each processing region..

1 20. The apparatus of claim 19, wherein each substrate stripping chamber is an

2 oxidation chamber.

1 21. The apparatus of claim 20, wherein the oxidation chamber and is connected to a

2 remote plasma system having a RF generator or a microwave generator.

The apparatus of claim 12, wherein the multi-slot pre-heating module is disposed within the loadlock chamber.

- 1 23. A process for forming a mesoporous oxide film on a substrate, comprising:
- a) forming a sol gel precursor comprising a silicon/oxygen compound, an
- organic solvent, water, and a surfactant;
- b) depositing the sol gel precursor on the substrate;
- c) curing the deposited sol gel precursor to form an oxide film; and
- d) exposing the film to an oxidizing environment to form a mesoporous oxide film.
- 1 24. The process of claim 23, wherein the mesoporous oxide film comprises a structure
- 2 of interconnected pores of uniform diameter.
- 1 25. The process of claim 24, wherein the mesoporous oxide film further comprises a
- 2 cubic phase structure.
- 1 26. The process of claim 23 wherein the silicon/oxygen compound precursor is
- 2 selected from the group consisting of tetraethylorthosililate, tetramethoxy silane,
- 3 phenyltriethyloxy silane, methyltriethoxy silane, and combinations thereof.
- 1 27. The process of claim 23, wherein the organic solvent is selected from the group
- 2 consisting of ethanol, isopropanol, n-propanol, n-butanol, sec-butanol, t-butanol, ethylene
- 3 glycol and combinations thereof.
- 1 28. The process of claim 23, wherein the surfactant is a non-ionic surfactant selected
- 2 from the group consisting of polyoxyethylene oxides-propylene oxide-polyethylene oxide
- 3 triblock copolymers, octaethylene glycol monodecyl ether, octaethylene glycol
- 4 monohexadecyl ether, and combinations thereof.
- 1 29. The process of claim 23, further comprising adding an acid or base catalyst to the

- 1 sol gel precursor prior to deposition of the sol gel precursor.
- 1 30. The process of claim 23, wherein the oxidizing environment is a plasma
- 2 comprising a reactive oxygen species.
- 1 31. The process of claim 30, wherein the reactive oxygen species is ozone.
- 1 32. The process of claim 23, wherein the oxidizing environment is maintained at a
- temperature between about 200°C to about 400°C.
- 1 33. The process of claim 23, wherein the oxide film is exposed to the oxidizing
- 2 environment for about 30 to about 300 seconds.
- 1 34. The process of claim 23, wherein the mesoporous oxide film exhibits a dielectric
- 2 constant between about 1.6 and about 2.2.
- 1 35. The process of claim 23, wherein the mesoporous oxide film has a porosity of at
- 2 least 50%.
- 1 36. The process of claim 23, wherein the mesoporous oxide film is cured at a
- 2 temperature between about 50°C to about 450°C.
- 1 37. The process of claim 34, wherein the mesoporous oxide film is cured between
- 2 about 1 minute to about 10 minutes.
- 1 38. The process of claim 23, further comprising silylating the mesoporous oxide film
- 2 to render the mesoporous oxide film hydrophobic.
- 1 39. The process of claim 38, wherein the silylating the mesoporous oxide film is
- 2 performed by a slylating agent selected from the group consisting of tetramethyl disilazane
- 3 (TMDS), hexamethyl disilazane (HMDS), dimethylaminotrimethyl silane, and

- 1 combinations thereof.
- 1 40. The process of claim 39, wherein the silylation process is performed at a
- 2 temperature between about 25°C to 200°C.
- 1 41. The process of claim 40, further comprising depositing a capping layer on the
- 2 mesoporous oxide film.
- 1 42. The process of claim 41, wherein the capping layer is comprised of materials
- 2 selected from the group consisting of silicon nitride, silicon dioxide, silicon oxynitride,
- 3 amorphous silicon carbide, and combinations thereof.
- 1 43. A process for forming a mesoporous oxide film on a substrate, comprising:
- a) introducing a substrate into a chamber;
- depositing a sol gel precursor on the substrate to form an oxide film, the sol
- 4 gel precursor comprising a silicon/oxygen compound, an organic solvent, water, and a
- 5 surfactant; and
- c) removing the organic solvent, water, and the surfactant from the oxide film
- by heating the film at a temperature of about 200°C to about 450°C in an inert atmosphere
- 8 to form a mesoporous oxide film.
- 1 44. The process of claim 43, wherein the mesoporous oxide film comprises a structure
- 2 of interconnected pores of uniform diameter.
- 1 45. The process of claim 44, wherein the mesoporous oxide film further comprises a
- 2 cubic phase structure.
- 1 46. The process of claim 43, wherein the silicon/oxygen compound precursor is
- 2 selected from the group consisting of tetraethylorthosililate, tetramethoxy silane,
- 3 phenyltriethyloxy silane, methyltriethoxy silane, and combinations thereof.

- 1 47. The process of claim 43, wherein the organic solvent is selected from the group
- 2 consisting of ethanol, isopropanol, n-propanol, n-butanol, sec-butanol, t-butanol, ethylene
- 3 glycol and combinations thereof.
- 1 48. The process of claim 43, wherein the surfactant is a non-ionic surfactant selected
- 2 from the group consisting of polyoxyethylene oxides-propylene oxide-polyethylene oxide
- 3 triblock copolymers, octaethylene glycol monodecyl ether, octaethylene glycol
- 4 monohexadecyl ether, and combinations thereof.
- 1 49. The process of claim 48, further comprising adding an acid or base catalyst to the
- 2 sol gel precursor prior to deposition of the sol gel precursor.
- 1 50. The process of claim 43, wherein the inert atmosphere comprises a non-reactive
- 2 gas selected from the group consisting of nitrogen, helium, argon, and combinations
- 3 thereof.
- 1 51. The process of claim 43, wherein the mesoporous oxide film is formed by
- annealing the oxide film at a temperature between about 400°C to about 450°C.
- 1 52. The process of claim 43, wherein the mesoporous oxide film is annealed for about
- 2 30 to about 300 seconds.
- 1 53. The process of claim 43, wherein the mesoporous oxide film exhibits a dielectric
- 2 constant between about 1.6 and about 2.2.
- 1 54. The process of claim 43, wherein the atmosphere comprises an oxidizing
- 2 environment of reactive oxygen species.
- 1 55. The process of claim 54, wherein the oxidizing environment is maintained at a
- 2 temperature between about 200°C to about 400°C.

- 1 56. The process of claim 55, wherein the oxide film is exposed to the oxidizing
- 2 environment for about 30 to about 300 seconds.
- 1 57. The process of claim 43, further comprising silylating the mesoporous oxide to
- 2 render the film hydrophobic.
- 1 58. The process of claim 43, wherein the silylation process is performed with a
- 2 slylating agent selected from the group consisting of tetra-methyl-di-silazane (TMDS),
- 3 hexamethyl disilazane (HMDS), deimethylaminotrimethyl silane, and combinations
- 4 thereof.
- 1 59. The process of claim 43, wherein the silylation process is performed at a
- 2 temperature between about 25°C to 200°C.
- 1 60. The process of claim 43, further comprising depositing a capping layer on the
- 2 mesoporous oxide film.
- 1 61. The process of claim 60, wherein the capping layer is comprised of materials
- 2 selected from the group consisting of silicon nitride, silicon dioxide, silicon oxynitride,
- 3 amorphous silicon carbide, and combinations thereof.

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